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(58) Field of search

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C3J

C3M

D1P

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C08L

(54) A thickening agent and cosmetic compositions containing it

(57) A gelling or thickening agent is produced from the ionic interaction of:

a cationic polymer comprising a polymer of a cellulose, or a cellulose derivative, which is grafted with a quaternary ammonium salt of a water-soluble monomer, and

a carboxylic anionic polymer having a specified capillary viscosity and Epprecht-Drage viscosity.

The anionic polymer may be polymethacrylic acid, a copolymer of methacrylic acid with an alkyl acrylate or methacrylate, an acrylamide derivative, maleic acid, a monoalkyl maleate or N-vinyl pyrrolidone, or an ethylene-maleic anhydride copolymer.

The agent is incorporated in compositions for treating the hair, skin or nails e.g. hair rinsing or setting lotions, shampoos, anti dandruff compositions, anti seborrhoeic compositions, support gels for permanent waving, hair dyeing compositions, anti-acne compositions and antipsoriatic compositions.

	A thickening agent and cosmetic compositions containing it	
5	The present invention relates to a new gelling or thickening agent, new thickened or gelled cosmetic compositions containing such an agent and a process enabling cosmetic compositions to be gelled and/or thickened.	. 5
10	A general requirement existing in the cosmetics industry is for compositions for hair or for the skin which do not flow too quickly; such is the case, in particular, with the compositions employed in processes which involve periods of application or of contact of the composition with the hair or the skin. It is very advantageous, in this case, to employ compositions which have a viscosity index higher than a certain limit enabling the products to be properly localized with the aid of thickened solutions.	10
15	In previous patents such as French Patents 2,383,660, 2,505,179 and 2,542,997, we have already described compositions containing cationic polymers and anionic polymers in an aqueous medium capable of being presented in the form of thickened or gelled compositions. The	15
20	resulting from the presence of the gelling or thickening agents, of excessively loading the hair or of leaving an unattractive powdery deposit or, yet again, of imparting to it an unpleasant feel or a dull appearance, particularly when involving compositions whose application is not followed by	20
25	are sometimes cloudy or opaque, and this can prevent their use in certain applications such as, for example, hair-shaping compositions which are generally clear.	25
30	We have investigated the possibility of preparing gelled or thickened aqueous cosmetic compositions conferring onto hair the advantageous shape-retention and sheen properties of the compositions containing cationic and anionic polymers, while avoiding the abovementioned disadvantages due to the addition of gelling agents or thickeners.	30
35	It is known to form gels from a polymer derived from a quaternary ammonium of cellulose ether as described in US-A-3,472,840 and from an anionic polymer which is alginic acid or a polysulphonic acid such as 2-acrylamido-2-methylpropanesulphonic acid. The gelled compositions produced in this manner result, on the one hand, from the use of anionic polymers which themselves have thickening or gelling properties and, furthermore, require relatively high solids concentrations. Furthermore, such compositions are not completely satisfactory when they are employed for conditioning hair damaged by physical or chemical treatments or by atmospheric	35
40	or thickened by a copolymer of cellulose or of a cellulose derivative which are grafted by a radical route with a quaternary ammonium salt of a water-soluble monomer with certain carboxy-	40
45	lic anionic polymers. This synergistic effect appears to be due, though this is merely a hypothesis, to the formation of an interpolymer by ionic interaction in an aqueous medium. To make the definition easier, the term "thickener" or "thickening agent" is employed in the remainder of the specification to denote a product having thickening and/or gelling properties resulting from this interaction.	45
50	The formation of a thickening agent is particularly surprising insofar as it results from polymers which do not individually have the thickening properties of the resulting agent. This capacity is markedly superior to that of gels known previously, some of which have been produced using anionic polymers which themselves have gelling properties. This is particularly advantageous within the scope of the present invention because the thickening characteristics make it possible not only to achieve a saving in the use of the polymers to obtain an identical gelling but at the	50
55	same time make it possible to impart to the hair or to the skin, which are treated with these compositions, certain improved cosmetic properties without loading the hair excessively. The cosmetic compositions containing the thickening agent have the advantage of not loading the hair, even when the applications are repeated, especially in the case of compositions which are applied using methods which do not involve a rinsing stage, and of imparting a pleasant feel	55
60	and a gleaming appearance to the hair. They impart good shape retention and good liveliness to hair, and more particularly to fine hair, in the case of the compositions whose application is	60

of damaged hair, especially insofar as its disentangling, its softness and its feel are concerned. The subject of the present invention concerns a thickener resulting from an ionic interaction in an aqueous medium of a copolymer of a cellulose or a cellulose derivative grafted by a radical 65 route with a quaternary ammonium salt of a water-soluble monomer with a particular group of

60 hair, and more particularly to fine hair, in the case of the compositions whose application is followed by a water rinse. Lastly, these compositions make it possible to improve the treatment

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.' '	carboxylic anionic polymers. The present invention provides a gelling or thickening agent produced from the ionic interac-	
	tion of: a cationic polymer comprising a polymer of cellulose or a cellulose derivative which are grafted with a quaternary ammonium salt of a water-soluble monomer, and a carboxylic anionic polymer having an absolute capillary viscosity, at a concentration of 5% in dimethylformamide or methanol at 30°C, of lower than or equal to 30×10 ⁻³ Pa s, this thickener having an Epprecht-Drage viscosity, module 3, of at least 0.45 Pa s in solution at a concentra-	5
10	preferably carried out by a radical route. The cationic polymer preferably has an absolute capillary viscosity at 1% in water at 30°C of less than 0.025 Pa s.	10
15	The cationic polymer is preferably a cellulose, or hydroxyalkyl cellulose such as hydroxymethyl cellulose, hydroxyethyl cellulose or hydroxypropyl cellulose which are grafted by a radical route with a methacryloylethyltrimethylammonium, methacrylamidopropyltrimethylammonium or dimethyldiallylammonium salt, more particularly a halide such as a chloride, or a methosulphate. A particularly preferred cationic polymer is a hydroxyethyl cellulose copolymer grafted by a	15
. 20	radical route with diallyldimethylammonium chloride sold under the trade name "Celquat L 200" or "Celquat H 100" by National Starch, which is also called "Polyquaternium 4" in the CFTA dictionary. When diluted to a concentration of 1% in water at a temperature of 30°C, this polymer has an absolute capillary viscosity of the order of 0.01 Pa s in the case of the product marketed under the trade name "Celquat L 200" or of 0.021 Pa s in the case of the product	20
2!	marketed under the trade name "Celquat H 100". The carboxylic anionic polymer preferably has a molecular weight of from 500 to 3,000,000 more particularly from 1,000 to 3,000,000. It is preferably a film-forming polymer. Particularly preferred polymers are:	25
	(a) a methacrylic acid homopolymer which has a molecular weight of greater than 20,000, as determined by light scattering.	
30	an acrylamide derivative, such as N,N-dimethylacrylamide, diacetoneacrylamide or N-tert-butyla-	30
3!	crylamide; maleic acid; C ₁ -C ₄ monoalkyl maleate; or N-vinylpyrrolidone; or (c) a copolymer of ethylene with maleic anhydride, such as the product sold under the trade name EMA 31 by Monsanto Cie.	35
41	Particularly preferred anionic polymers are methacrylic acid copolymers which have an absolute capillary viscosity measured at a concentration of 5% in solution in dimethylformamide or methanol, at 30°C, of from 0.003 to 0.030 Pa s, more particularly a copolymer of methacrylic acid with methyl methacrylate whose absolute capillary viscosity, measured at a concentration of 5% in solution in dimethylformamide, is of the order of 0.015 Pa s or a copolymer of metha-	40
4	crylic acid with monoethyl maleate which has an absolute capillary viscosity, measured at a concentration of 5% in solution in dimethylformamide, of the order of 0.013 Pa s, a copolymer of methacrylic acid with butyl methacrylate whose absolute capillary viscosity, measured at a concentration of 5% in solution in methanol, is of the order of 0.010 Pa s, or a copolymer of methacrylic acid with maleic acid whose absolute capillary viscosity, measured at a concentration of 5% in solution in dimethylformamide, is of the order of 0.016 Pa s.	45
5	The thickener may, for example, be prepared under the following conditions: a quantity of water is added to the copolymer of cellulose or cellulose derivative grafted by a radical route with a quaternary ammonium salt of a water-soluble monomer to dissolve it	50
5	(solution 1). Separately, a quantity of water is added to the carboxylic anionic polymer to dissolve it, the dissolution being promoted by neutralization with a conventional alkalifying agent such as aqueous ammonia or an alkanolamine (solution II). The thickener may then be formed by adding solution I to solution II or vice versa, with stirring, at ambient temperature. When the gelling or thickening agent has formed it can then, if	55
6	desired, be diluted with water or with a mixture of water and alcohol, the proportion of alcohol being that required to produce the required alcoholic strength for the formulation. O According to an alternative form of this process, it is equally possible, without recourse to neutralization, to dissolve the carboxylic anionic polymer in alcohol, preferably ethanol, at a	60
	concentration such as to bring the final formulation to the alcoholic strength required. The thickener may also be formed in the aqueous cosmetic medium itself. The copolymer of cellulose or a cellulose derivative which are grafted with a quaternary	

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to 6%, especially 0.1 to 1.5%, by weight relative to the weight of the composition. The carboxylic anionic polymer is preferably used in an aqueous medium, generally in an amount of from 0.01 to 6%, especially 0.1 to 1.5%, by weight relative to the weight of the composition. The Weight ratio of the cationic polymer to the carboxylic anionic polymer is preferably from 1:5 to 5:1, more preferably from 1:2 to 2:1 and is especially equal to about 1:1. The present invention also provides a cosmetic composition suitable for the treatment of hair, skin or nails which comprises at least one gelling or thickening agent as defined above and at least one further adjuvant. The thickener is preferably present in the composition of the present invention in a concentra-10 tion of from 0.02 to 12%, more preferably from 0.2 to 3%, by weight based on the total 10 weight of the composition. This composition is generally in aqueous form, but may contain other cosmetically acceptable solvents such as, for example, lower (for example C1-C8 or C1-C4) alcohols such as ethanol or isopropanol, glycerol, glycols or glycol ethers such as ethylene glycol monobutyl ether, propylene 15 glycol, diethylene glycol monoethyl ether and monomethyl ether, in proportions which do not 15 affect the formation of the thickener. These compositions have a pH which is generally from 6 to 12, preferably from 6.5 to 9, more particularly, close to neutrality, for example of the order of 7 to 8. The pH may be adjusted with an alkalifying or acidifying agent which is usually employed in 20 the field of cosmetics. 20 The cosmetic composition may, for example, be employed as a shampoo, after-shampoo composition, product for rinsing to be applied before or after shampooing, before or after dyeing or bleaching, before or after permanent-waving or hair straightening, a hair-setting or blow-drying composition, a restructuring composition, or a support for permanent-waving or for dyeing or 25 25 bleaching hair. The composition may also contain a dermatological active principle such as an antidandruff, antiseborrhoeic, antiacne, antifungal, bactericidal, keratolytic or antipsoriatic agent. When the composition is in the form of a thickened lotion or gel for hair-setting or for blowdrying, it may optionally contain other polymers which are usually employed in a composition of this type, more particularly nonionic polymers such as polyvinylpyrrolidones, copolymers of 30 polyvinylpyrrolidone with vinyl acetate, or anionic polymers which do not have the abovemen-30 tioned properties of gelling or thickening with the cationic polymer, for example copolymers of vinyl acetate with an unsaturated carboxylic acid such as crotonic acid, copolymers resulting from the copolymerization of vinyl acetate with crotonic acid and an acrylic or methacrylic ester, copolymers resulting from the copolymerization of vinyl acetate with an alkyl vinyl ether and an 35 35 unsaturated carboxylic acid and copolymers resulting from the copolymerization of vinyl acetate with crotonic acid and a vinyl ester of an acid containing a long carbon chain or an allyl or methallyl ester of an acid containing a long carbon chain. These polymers are generally employed in a concentration of from 0.1 to 5% by weight based on the total weight of the composition. 40 When employed as a rinsing composition, the composition may contain various conditioning agents such as quaternary proteins, cationic silicone polymers, cationic surfactants and cationic polymers other than polymers of cellulose or of cellulose derivatives grafted by a radical route with a quaternary ammonium water-soluble monomer, of the polyamine, polyaminoamide or quaternary polyammonium type. When the compositions are employed as shampoos, they may contain surface-active agents with detergent properties which are known per se, such as anionic, cationic, nonionic or amphoteric surface-active agents or mixtures thereof. In general, the surface-active agents are present in a proportion of from 0.1 to 30% by weight based on the total weight of the composition. 50 When the composition is employed for dyeing hair, it may contain a direct dye or oxidation dve precursor which is known in the art. The compositions may also be used for conditioning skin and nails.

ide or methanol at a concentration of 5%, is from 0.010 to 0.015 Pa s, the Epprecht-Drage 60 viscosity of the thickener, measured at 21°C, module 3, diluted to a concentration of 1% in water, being higher than 0.45 Pa s, and the pH of the composition being from 6.5 to 9.

The compositions according to the invention may contain any other ingredient which is usually employed in cosmetics, such as perfumes, colourants, preservatives, sequestering agents, softeners or silicones.

A particularly perferred cosmetic composition is a hair-shaping composition which is not rinsed off. This composition comprises, in an aqueous or aqueous-alcoholic medium, a thickener result-

55 ing from the ionic interaction of 0.1 to 1.5% by weight of a hydroxyethyl cellulose copolymer grafted by a radical route with diallyldimethylammonium chloride and 0.1 to 1.5% by weight of a copolymer of methacrylic acid with methylmethacrylate or with monoethyl maleate or with butyl methacrylate whose absolute capillary viscosity, measured at 30°C in solution in dimethylformam-

The present invention also provides a process for thickening or gelling a cosmetic composition

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wherein at least one thickener as defined above or a composition containing the polymers forming the thickener in a proportion of from 0.02 to 12% by weight based on the total weight of the composition is introduced into the composition to give it an Epprecht-Drage viscosity, measured at 21°C (module 3), of at least 0.450 Pa s.

Aqueous gels or thickened compositions containing the thickener may be prepared separately, and the cosmetic composition may be prepared in a different step, if desired at the time of use.

The present invention also provides a process for the treatment of hair, of the skin and of the nails, wherein a cosmetic composition as defined above is applied thereto, it being possible for this composition to be rinsed off with water, or not, according to the nature of the treatment 10 desired.

We have found that the composition for the treatment of hair not only makes it possible to localize the product on hair properly without flowing onto the face but that the hair treated in this manner also has a pleasant feel and a shiny appearance. Furthermore, the thickened or gelled composition has the advantage of being clear.

15 The examples which follow further illustrate the invention.

EXAMPLE 1

Aqueous gels were prepared according to the information which appears in Table A which follows. For this purpose 50 cm³ of an aqueous solution containing 1% of active substance of the product marketed under the trade name of "Celquat L 200", which is a copolymer of hydroxyethyl cellulose grafted by a radical route with diallyldimethylammonium chloride, were added at ambient temperature and with mechanical stirring to 50 cm³ of an ethanolic solution at an alcohol strength of 20° containing 1% as active substance of the previously neutralized anionic polymer defined in the table.

In Table A below, the measurement of the absolute capillary viscosity of the anionic polymers is carried out in dimethylformamide (DMF) and/or in methanol.

45

TABLE A

CATIONIC POLYMER				Epprecht-Drage viscosity	
The state of the s		Absolute capillary viscosity Pa s x 10 ⁻³		of the thickener formed Pa s	
CELQUAT L 200		(1) 10.	١.		
CARBOXYLIC ANIONIC POLYMER	Propor-	(2) DHP	CE 30E		
Methacrylic acid/methyl, methacrylate copolymer	50/50	15	 	1-550	
	80/20	24-47	10.56	1- 430	
Methacrylic acid/methyl acrylate copolymer	50/50		16-4	1, 300	
	80/20	17.7	8-5	1. 150	
Methadrylic acid/butyl methacrylate copolymer	65/15		9-94	2,000	
Methacrylic acid/monoethyl melente copolymer	63-6/ 36-4	3,46		0.620 (mod 4)	
	59/41	8	}	1-000 (mod 4)	
	66/34	19-2		0,780;1,500 (mod 4)	
	61/39	26.8		0-580;1-250 (mod 4)	
	62/38	10-4		0.550;1.000 (mod 4)	
	65/35	14,1		0.600/1.200 (mod 4)	
n	66/34	13 12		1,490;2,000 (mod 4) 1,700;2,100 (mod 4)	
	68/32	19.2		1.700;2.500 (mod 4)	
W W	72/2B	14.2		1, 380 ; 1, 500 (mod 4)	
Methacrylicacid/N,N-dimethylacrylemide copolymer	50/50			0.980	
•	80/20	16-3 -		1.350	
Methacrylic acid/diacetoneacrylamide copolymer (4)	80/20		1,07	1-200	
Methacrylic acid/N-tert-butylacrylamide copolymer	80/20		4.06	1-050	
Methacrylic acid/maleic acid copolymer	65/35	16.7		2-100	
Methacrylic acid/N-vinylpyrrolidone copolymer _	70/30	13.6		1-800	
	80/20	9-2		1-050	
Polymethacrylic acid MI 137,000			6.8	1.400	
" " MM 186,000	1		9,8	2.100	

⁽¹⁾ measured at 30°C in 1% strength solution in water

⁽²⁾ measured at 30°C in 5% strength solution in dimethylformamide or methanol

⁽³⁾ module 3 - measured at 21° C in 1% strength 10° aqueous sicohol solution - pH = 7.5

⁽⁴⁾ viscosity measured using a 1% atrength solution of this amionic polymer.

EXAMPLES 2 to 11

The following gelled compositions for hair styling are prepared (Tables B and C).

When these various compositions are applied to clean wet hair, they impart shape retention to it without leaving a powdery deposit. When they are applied to dried hair it is found that the composition makes styling easier without loading the hair and that, once dried, the latter is soft and has a pleasant feel.

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		EXAMPLE N	PLE No.			
COMPOSITIONS	2	3	4	5	6]
Celcust H 100 g X AS	0.5	0.4				
Celquat L 200 g Z AS			0-8	1	0-3	
Methacrylic acid/monoethyl maleste copolymer (66/34) g % AS	0.5			·		ŀ
Methacrylic acid/maleic acid copolymer (70/30) g % AS		0.6		'		
Rethacrylic acid/butyl methacrylate copolymer (85/15) g % AS			0.8		·	
Polymethacrylic acid Mw 137,000 g X AS	. ,	' '			0.4	١, .
Ethylene/maleic anhydride copolymer Monsanto ENA 31 g % AS	,			0.8		
: 2-Amino-2-methyl-1-propanol q.s. pH		9	,	6	9	
Ethyl alcohol q.a.	20"		25-		10	
Nater q.s. g	100	100	100	100	100	
Epprecht-Orage viscosity 21 ⁰ C 1% in H ₂ O (module 3) in Pa s	L 150	0, 700	2 -150	2,400	0 .725	

TABLE C

5							5
		EXAMPLE No.				1	
•	COMPOSITIONS	7	8	9	10	11	
10	Celcust H 100 g X AS	0.4			0.5		10
	Celquat L 200	l	1	0.66		0-33	
15	Nethacrylic acid/M-tert-butylacrylamide copolymer 80/20 g % AS	0.2					4=
15	Rethacrylic acid/N_H-dimethyl acrylamide copolymer 80/20 g % AS		0-5			į	15
	Rethacrylic acid/methyl methacrylate copolymer 50/50 g % AS			0.33			
20	Methacrylic scid/methyl methacrylate copolymer : 80/20 g % AS				1.		20
	Polymethacrylic acid RM 186,000 g % AS					0.66	
				!			• • •
25	2-As ino-2-sethy l-1-propanol					,	25
	q.s. pil		8.5	7.5	8-5	7.5	
	Ethyl alcohol		30°	10° .	10°	10°	
30	Perfuse, colorant, preservative Water						30
	q.s. g Epprecht-brage viscosity 21°C	100	100	100	100	100	
	1% in H ₂ O (module 3) in Pa s	0.480	1.800	0, 900	1. 725	1. 300	
35	-			•			35
40	EXAMPLE 12 An after-shampoo of the following composition is pre (A) Celquat L 200 from National Starch O. (B) 72/28 Methacrylic acid/monoethyl maleate copolymer O.	7 g A	\s	,			40
40	copolymer Distearyldimethylammonium chloride Hydrochloric acid q.s. pH; 7 Water q.s. 100	7 g <i>f</i> 9 g	45				40
45	This composition is applied to clean, roughly dried had minutes it is rinsed off with water. The wet hair is smooth and has body. The gel obtained by interaction of the two polymers	ir. Afte	d slippe	ery. Afte	r drying i	it is lively	
50	at 21°C, module 3, of 1.7 Pa s at a concentration of 1	.4% in	water.	an Eppro	one broge		
50	EXAMPLE 13 An after-shampoo of the following composition is pre (A) Celquat L 200 from National Starch O.	<u>.</u> .					50
EE	(B) 50/50 Methacrylic acid/methyl methacrylate						FF
55	Quaternized protein sold under the trade	•				.*	55
	name of "Lexein QX 3000" by Inolex 1. Hydrochloric acid q.s. pH: 6.7	g A	NS.				
60	Water q.s. 100	9		:			60
	This gelled composition is applied to clean, roughly d few minutes it is rinsed off with water. The dried hair is lively and has body. The gel obtained by interaction of the network A composition of the polymers.			-			
65	The gel obtained by interaction of the polymers A an 21°C, module 3, of 1.8 Pa s at a concentration of 1.49	in wa	s an Ep ater.	precnt-D	rage visc	osity at	65

	EXAMPLE 14			•		
	The following shampoo is prepared:				•	
•	(A) Celquat L 200 from National Starch	0.5	g AS			, -
5	(B) 50/50 Methacrylic acid/methyl methac	rylate 0.7	g AS		•	' 5
	copolymer Nonionic surfactant of formula:	0.7	g AS			<i>t</i> ,
	Monitorite surfactant or formula.					
	R-CHOH-CH2O-[CH2-CHOH-CH2O],-H					10
10						. 10
	in which R denotes a mixture of C ₉ -C ₁₂ alkyl radic	als .		•		
	n denotes a statistical mean value of			·	•	
	about 3.5	10	g AS		•	. 45
15	Hydrochloric acid q.s. pH: 7.	4 ·				15
	Perfume, preservative q.s. Water	100	g	•		
	AAGIGI		9			
	This shampoo has the appearance of a c	ear gel.				20
20	The gel obtained by interaction of the po	lymers A and	B has an	Epprecht-D	rage viscosity	y at 20
	21°C, module 3, of 1.65 Pa s at a concent	ration of 1961	n water.			-
	EXAMPLE 15					
	The following shampoo is prepared:					25
25	(A) Celquat L 200 from National Starch	0.7	g AS			25
	(B) 72/28 Methacrylic acid/monoethyl mal copolymer	0.7	g AS			
	Sodium alkyl ether carboxylate oxyethyler		3			
	with 3 moles of ethylene oxide, sold by				•	20
30	Marchon under the trade name "Empilan	· 10	- 40			30
	2747/30" Hydrochloric acid q.s. pH: 6	10	g AS			
	Perfume, preservative q.s.					
	Perfume, preservative q.s. Water q.s.	1Ó0	9			. 25
35	Water q.s.		g			35
35	Water q.s. This shampoo has the appearance of a c	lear gel.		ı Epprecht-D	rage viscosit	
35	Water q.s.	lear gel. lymers A and	B has an	Epprecht-D	rage viscosit	
	Water q.s. This shampoo has the appearance of a c The gel obtained by interaction of the po 21°, module 3, of 1.7 Pa s at a concentrat	lear gel. lymers A and	B has an	Epprecht-D	rage viscosit	y at
	Water q.s. This shampoo has the appearance of a c The gel obtained by interaction of the po 21°, module 3, of 1.7 Pa s at a concentrat EXAMPLE 16	lear gel. lymers A and	B has an	Epprecht-C	Prage viscosit	
	Water q.s. This shampoo has the appearance of a c The gel obtained by interaction of the po 21°, module 3, of 1.7 Pa s at a concentrat EXAMPLE 16 The following lotion is prepared:	lear gel. lymers A and	B has an water.	Epprecht-C	rage viscosit	y at
	Water q.s. This shampoo has the appearance of a c The gel obtained by interaction of the po 21°, module 3, of 1.7 Pa s at a concentrat EXAMPLE 16 The following lotion is prepared: (A) Celquat L 200 (B) Polymethacrylic acid	lear gel. lymers A and ion of 1.4% ir 0.1	B has an	n Epprecht-D	rage viscosit	y at
40	Water q.s. This shampoo has the appearance of a c The gel obtained by interaction of the po 21°, module 3, of 1.7 Pa s at a concentrat EXAMPLE 16 The following lotion is prepared: (A) Celquat L 200 (B) Polymethacrylic acid 2-Amino-2-methyl-1-propanol q.s. pH: 7.	lear gel. lymers A and ion of 1.4% ir 0.1	B has an water.	a Epprecht-D	rage viscosit	y at 40
	Water q.s. This shampoo has the appearance of a c. The gel obtained by interaction of the po 21°, module 3, of 1.7 Pa s at a concentrate EXAMPLE 16 The following lotion is prepared: (A) Celquat L 200 (B) Polymethacrylic acid 2-Amino-2-methyl-1-propanol q.s. pH: 7. Perfume, colorant, preservative q.s.	lear gel. lymers A and ion of 1.4% ir 0.1 0.1	B has an water.	Epprecht-C	rage viscosit	y at
40	Water q.s. This shampoo has the appearance of a c The gel obtained by interaction of the po 21°, module 3, of 1.7 Pa s at a concentrat EXAMPLE 16 The following lotion is prepared: (A) Celquat L 200 (B) Polymethacrylic acid 2-Amino-2-methyl-1-propanol q.s. pH: 7.	lear gel. lymers A and ion of 1.4% ir 0.1	B has an water.	Epprecht-C	rage viscosit	y at 40
40	Water q.s. This shampoo has the appearance of a c. The gel obtained by interaction of the po 21°, module 3, of 1.7 Pa s at a concentrate EXAMPLE 16 The following lotion is prepared: (A) Celquat L 200 (B) Polymethacrylic acid 2-Amino-2-methyl-1-propanol q.s. pH: 7. Perfume, colorant, preservative q.s. Water q.s. This hair-setting lotion is slightly gelled a	lear gel. lymers A and ion of 1.4% ir 0.1 0.1 5 100 nd does not re	B has an water. 9 9 9	sing.		y at 40 45
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40 45	This shampoo has the appearance of a compact the gel obtained by interaction of the post of the gel obtained by interaction of the post of the gel obtained by interaction of the post of the gel obtained by interaction of the post of the gel obtained by interaction of the post of the gel obtained by interaction of the post of the gel obtained by interaction of the post of the gel obtained by interaction of the post of the gel obtained by interaction of the post of the gel obtained by interaction of the post of the gel obtained by interaction of the post of the gel obtained by interaction of the post of the gel obtained by interaction of the post of the gel obtained by interaction of the gel obtained by inter	lear gel. lymers A and ion of 1.4% ir 0.1 0.1 5 100 and does not relymers A and antration of 0.2 prepared: 1.5	B has an water. 9 9 9 equire rin B has ar % in wat	sing. 1 Epprecht-L		y at 40 45 y at
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40 45	This shampoo has the appearance of a contract of the gel obtained by interaction of the post of the gel obtained by interaction of the post of the gel obtained by interaction of the post of the following lotion is prepared: (A) Celquat L 200 (B) Polymethacrylic acid 2-Amino-2-methyl-1-propanol q.s. pH: 7. Perfume, colorant, preservative q.s. Water q.s. This hair-setting lotion is slightly gelled a The gel obtained by interaction of the post of the gel obtained by interaction of the post o	lear gel. lymers A and ion of 1.4% ir 0.1 0.1 5 100 Ind does not relymers A and intration of 0.2 In prepared: 1.5 leate 1.2 tyl)- l under	B has an water. 9 9 equire rin B has ar % in wat	sing. 1 Epprecht-L		y at 40 45 y at 50
40 45 50	This shampoo has the appearance of a contract of the gel obtained by interaction of the post of the gel obtained by interaction of the post of the gel obtained by interaction of the post of the following lotion is prepared: (A) Celquat L 200 (B) Polymethacrylic acid 2-Amino-2-methyl-1-propanol q.s. pH: 7. Perfume, colorant, preservative q.s. Water q.s. This hair-setting lotion is slightly gelled a The gel obtained by interaction of the post of the gel obtained by interaction of the post o	lear gel. lymers A and ion of 1.4% ir 0.1 0.1 5 100 and does not relymers A and antration of 0.2 prepared: 1.5 leate 1.2 tyl)-	B has an water. 9 9 9 equire rin B has ar % in wat	sing. 1 Epprecht-L		y at 40 45 y at 50 55
40 45	This shampoo has the appearance of a case of the gel obtained by interaction of the post of the post of the following lotion is prepared: (A) Celquat L 200 (B) Polymethacrylic acid 2-Amino-2-methyl-1-propanol q.s. pH: 7. Perfume, colorant, preservative q.s. Water q.s. This hair-setting lotion is slightly gelled a The gel obtained by interaction of the post of t	lear gel. lymers A and ion of 1.4% ir 0.1 0.1 5 100 Ind does not relymers A and intration of 0.2 In prepared: 1.5 leate 1.2 tyl)- l under	B has an water. 9 9 equire rin B has ar % in wat	sing. 1 Epprecht-L		y at 40 45 y at 50
40 45 50	Water q.s. This shampoo has the appearance of a c. The gel obtained by interaction of the po 21°, module 3, of 1.7 Pa s at a concentrate. EXAMPLE 16 The following lotion is prepared: (A) Celquat L 200 (B) Polymethacrylic acid 2-Amino-2-methyl-1-propanol q.s. pH: 7. Perfume, colorant, preservative q.s. Water q.s. This hair-setting lotion is slightly gelled a The gel obtained by interaction of the po 21°C, module 2, of 0.095 Pa s at a concentrate. EXAMPLE 17 The following antidandruff composition is (A) Celquat L 200 (B) 66/34 Methacrylic acid/monoethyl mail copolymer 1-Hydroxy-4-methyl-6-(2,4,4-trimethylpen 2-(1H)-pyridinone, ethanolamine salt, sold the trade name "Octopirox" by Hoechst Ethyl alcohol q.s. 30° 2-Amino-2-methyl-1-propanol q.s. pH 7	lear gel. lymers A and ion of 1.4% ir 0.1 0.1 5 100 Ind does not relymers A and intration of 0.2 In prepared: 1.5 leate 1.2 tyl)- l under	B has an water. 9 9 equire rin B has ar % in wat	sing. 1 Epprecht-L		y at 40 45 y at 50 55
40 45 50	This shampoo has the appearance of a contract of the gel obtained by interaction of the position of the position of the position of the position of the gel obtained by interaction of the position of the gel obtained by interaction of the position of the gel obtained by interaction of the position of t	lear gel. lymers A and ion of 1.4% ir 0.1 0.1 5 100 Ind does not relymers A and intration of 0.2 In prepared: 1.5 leate 1.2 tyl)- l under	B has an water. 9 9 equire rin B has ar % in wat	sing. 1 Epprecht-L		y at 40 45 y at 50 55
40 45 50	This shampoo has the appearance of a case of the gel obtained by interaction of the post of the gel obtained by interaction of the post of the following lotion is prepared: (A) Celquat L 200 (B) Polymethacrylic acid 2-Amino-2-methyl-1-propanol q.s. pH: 7. Perfume, colorant, preservative q.s. Water q.s. This hair-setting lotion is slightly gelled a The gel obtained by interaction of the post of th	lear gel. lymers A and ion of 1.4% ir 0.1 0.1 5 100 Ind does not relymers A and intration of 0.2 prepared: 1.5 leate 1.2 tyl)- 1 under 0.1	B has an water. g g g equire rin B has ar % in wat g g	sing. Epprecht-E ter	Orage viscosit	y at 40 45 y at 50 55

	The gel obtained by interaction of the polymers A and B 21°C, module 3, of about 1.8 Pa s at a concentration of 2.	has an Epprecht-Drage viscosity at 7% in water.
5	EXAMPLE 18 The following antiseborrhoeic composition is prepared: (A) Celquat L 200 0.5 g (B) 50/50 Methacrylic acid/methyl methacrylate	5
10	copolymer 0.5 g Poly- β -alanine 1 g	10
15	This antiseborrhoeic composition which can be applied to ance of a clear gel and does not require rinsing. The gel obtained by interaction of the polymers A and B I 21°C, module 3, of about 1.2 Pa s at a concentration of 1%	the skin or to hair has the appear-
20	Composition 1	emposition, is prepared: 20
25	Glycerol monothioglycolate 68.3 g Glycerin q.s. 100 g Composition 2 Celquat L 200 1.8 g	٥٠
	70/30 Methacrylic acid/maleic acid copolymer 1.5 g 2-Amino-2-methyl-1-propanol q.s. pH 6.5	25
30	Triethanolamine 3 g Perfume, colorant, preservative q.s. Water q.s. 100 g	30
35	The two compositions 1 and 2 are mixed ad hoc in propose 87 g of composition 2. This mixture is applied to hair which is wound onto rollers, in place, it is rinsed off and an oxidizing solution consisting c 3, is applied for 10 minutes. The hair is then rinsed.	for 15 minutes. After 15 minutes. 35
40	The following direct-dyeing composition is prepared:	40
45	Celquat L 200 from National Starch 1-N-(y-hydroxypropyl)amino-2-nitro-4-N',N'- bis(y-hydroxyethyl)aminobenzene monohydro-	AS AS 45
50	2-Amino-2-methyl-1-propanol q.s. pH 7.5 Ethyl alcohol q.s. 10°	50

This dyeing composition is applied to wet brown hair, washed beforehand. After drying, the hair acquires an ashen brown color.

	1		·					
	EXAMPLE 21			•	•			
	The antipsorlatic composition		ding 0.	5 g of	anthra	line at th	e time of use to	
5	the gel of the following composition (A) Celquet L 200	sition:	0.5	~ ,			•	5
. 5	(B) 50/50 Methacrylic acid/me	ethyl methacrylate	0.5	g				. 0
	copolymer	only. Monitor yidlo	0.5	g			1	
	2-amino-2-methyl-1-propanol	q.s. pH 7	1	•				
	Ethyl alcohol	q.s. 10°		٠,				
10	Preservative	q.s.						10
•	Water	q.s.	100	g				
15	The antipsoriatic composition The gel obtained by interactic 21°C, module 3, of about 1.2 F	on of the polymers	A and	B has	an Epp	precht-Dra	ing. age viscosity at	15
20	EXAMPLE 22 The following antiacne composition is applied to	sition is given in E			g of be	nzoyl per	oxide at the time	20
			1		٠,		•	
	EXAMPLE 23						•	
25	The following bactericidal con phenoxy)phenol or triclosan (DC to the gel whose composition is This composition is applied to	l) sold under the na s given in Example	ame of	dding ''Irgas	1 g of san DP	5-chloro- 300" at	2-(2,4-dichloro- the time of use	25
	EXAMPLE 24				•			
	A hair-conditioning composition	on is prepared by a	adding '	8 g o	f iris p	owder dil	uted with 36 g	•
30	of water to 46 g of a gel of the	e following compos	sition:	1	-		,	30
	(A) Celquat L 200		4.5	g			•	
	(B) 80/20 Methacrylic acid/N-v	rinylpyrrolidone		_	•	•		
	copolymer Ethyl alcohol	q.s. 10°	4.5	g		*	•	
35	2-Amino-2-methyl-1-propanol					•		35
-	Perfume, preservative	q.s.				•	•	.00
	Water	q.s.	100	.g			•	
40	The composition is applied to The gel obtained by interactio 21°C, module 4, of 11.7 Pa s a	n of the polymers	A and	B has	an Epp	s a soft i recht-Dra	eel. ge viscosity at	40
	EXAMPLE 25		•					
45	The following restructuring rin ourea at the time of use to the This composition is applied to	gel of Example 21	ared by at pH	adding 6.	g 1.5 g	of dime	thylolethylenethi-	45
	CLAIMS							
50	A gelling or thickening age a cationic polymer comprising grafted with a quaternary ammo	a polymer of a ce	flulose (or a ce	ellulose	derivativ	e which are	50
	a carboxylic anionic polymer h				•		ntration of 5% in	
. 56	dimethylformamide or methanol having an Epprecht-Drage viscos tion of 1% in water at 21°C.	at 30°C, of lower	than or	equal	to 30>	< 10 ⁻³ Pa	s, this thickener	55
	2. An agent according to cla	im 1 wherein the d	cationic	polvm	er is a	hydroxya	ikvi cellulose	55
	copolymer grafted by a radical r monomer which is a methacrylo ium or dimethyldiallylarmonium	oute with a quateri ylethyltrimethylamn	nary arr	moniu	m salt	of a wat	er-soluble	
60	 An agent according to cla a methacrylic acid homopolym 	im 1 or 2, wherein						60
65	by light scattering, a copolymer of methacrylic ac derivative, maleic acid, a C ₁ -C ₄ i a copolymer of ethylene with	monoalkyl maleate					n acrylamide	65
	a copolymor or employed with	maiero arirryantae.						-

		4. An agent according to any one of claims 1 to 3 wherein the anionic polymer is: a copolymer of methacrylic acid with methyl methacrylate whose absolute capillary viscosity, measured in solution in dimethylformamide at a concentration of 5% at 30°C, is of the order of 15×10 ⁻³ Pa s,	
	5	a copolymer of methacrylic acid with monoethyl maleate having an absolute capillary viscosity, measured in solution in dimethylformamide at a concentration of 5% at 30°C, of the order of 13×10^{-3} Pa s,	5
	10	a copolymer of methacrylic acid with butyl methacrylate whose absolute capillary viscosity, measured in solution in methanol at a concentration of 5% at 30°C, is of the order of 10×10 ⁻³ Pa s, or	10
		a copolymer of methacrylic acid with maleic acid whose absolute capillary viscosity, measured in solution in dimethylformamide at a concentration of 5% at 30°C, is of the order of 16×10 ⁻¹ Pa s. 5. An agent according to any one of claims 1 to 4 wherein the weight ratio of the cationic	
	15	6. An agent according to any one of claims 1 to 5 which has been prepared in an aqueous medium comprising 0.01 to 6% of the cationic polymer and 0.01 to 6% of the carboxylic anionic polymer.	15
	20	7. An agent according to claim 1 substantially as hereinbefore described with reference to any one of the Examples. 8. A cosmetic composition suitable for the treatment of hair, skin or nails, which comprises	20
		at least one gelling or thickening agent as defined in any one of claims 1 to 7 and at least one further adjuvant.	
•	25	9. A composition according to claim 8 wherein the gelling or thickening agent is present in a proportion of from 0.02 to 12% by weight based on the total weight of the composition. 10. A composition according to claim 8 or 9, which has a pH of from 6 to 12.	25
	30	11. A composition according to any one of claims 8 to 10 suitable for use as a thickened or gelled lotion for hair-setting or for blow-drying which additionally comprises a nonionic polymer which is a polyvinylpyrrolidone or copolymer or polyvinylpyrrolidone with vinyl acetate, or an anionic polymer which is a copolymer of vinyl acetate with an unsaturated carboxylic acid, a	30
		copolymer resulting from the polymerization of vinyl acetate with crotonic acid and an acrylic or methacrylic ester, a copolymer resulting from the copolymerization of vinyl acetate with a vinyl alkyl ether and an unsaturated carboxylic acid, a copolymer resulting from the copolymerization	
	35	of vinyl acetate with crotonic acid and a vinyl ester of an acid containing a long carbon chain or an allyl or methallyl ester of an acid containing a long carbon chain. 12. A composition according to any one of claims 8 to 11 in the form of a shampoo which comprises one or more anionic, cationic, nonionic or amphoteric surface-active agents with a	35
	40	detergent property. 13. A composition according to any one of claims 8 to 10, suitable for rinsing off, which	
	40	comprises a conditioning agent which is a quaternary protein, cationic silicone polymer, cationic surfactant or cationic polymer other than a polymer of a cellulose or cellulose derivative grafted by a radical route with a quaternary ammonium water-soluble monomer.	40
	45	14. A cosmetic composition suitable for use in hair-setting, which comprises, in an aqueous or aqueous-alcoholic medium, a thickener resulting from the ionic interaction of 0.1 to 1.5% by weight of a hydroxyethyl cellulose copolymer grafted by a radical route with diallyldimethylam-	45
	-	monium chloride and 0.1 to 1.5% by weight of a copolymer of methacrylic acid with methyl methacrylate or with monoethyl maleate or with butyl methacrylate whose absolute capillary viscosity, measured at 30°C in solution in dimethylformamide or methanol at a concentration of	•
	50	5%, is from 0.010 to 0.015 Pa s, the Epprecht-Drage viscosity of the thickener, measured at 21°C, module 3, diluted to a concentration of 1% in water, being higher than 0.45 Pa s, and the pH of the composition being from 6.5 to 9.	50
		 15. A cosmetic composition according to claim 8 or 14 substantially as hereinbefore described with reference to any one of the Examples. 16. A process for thickening or gelling an aqueous cosmetic composition wherein at least 	
	55	one thickener as defined in any one of claims 1 to 7 is introduced into the composition to give it an Epprecht-Drage viscosity measured at 21°C (module 3) of at least 0.45 Pa s at a concentration of 1% in water.	55
	60	 17. A process for the treatment of hair, of the skin or of the nails, wherein at least one cosmetic composition as defined in any one of claims 8 to 15 or produced by a process as defined in claim 16 is applied thereto. 18. A process according to claim 17 wherein a composition as defined in claim 11 or 14 is 	60
		applied, this application not being followed by a rinse.	